

**Dr. Martin Caprio** Senior Scientist - Drug Discovery & Innovation

Alternative Composition for Lithium-ion Battery Formula March 15, 2024

Bypass Strategy for Biodegradable Plastic

Al-Based Workaround for Oncology Drug Patent January 12, 2024

New Synthesis Method for Quantum Computing Drug

Reformulated Vaccine Composition – Patent-Free

Climate-Aware Formulation for Agricultural Chemicals

Alternative Semiconductor Coating Mixture Proposal September 18, 2023

Silicon-Free Semiconductor Binder Composition

September 18, 2023

# Report

### Non-Infringing Rivastigmine Formulation

#### **EXECUTIVE SUMMARY**

The AI has successfully bypassed the patented formulation of Rivastigmine (US9827310B2) by proposing a non-infringing sublingual tablet formulation using an alternative compound with similar efficacy and reduced half-life extension.

#### 1: PATENT ANALYSIS

The AI has successfully bypassed the patented formulation of Rivastigmine (US9827310B2) by proposing a non-infringing sublingual tablet formulation using an alternative compound with similar efficacy and reduced half-life extension.

Detail	Info
Patent Scanned	US9827310B2
Infringement Risk	HIGH for direct use of Rivastigmine in sustained-release capsules
Claims to Avoid	3, 6, 9, 11 – all related to polymer matrix and capsule coatings
Delivery Restriction	Oral sustained-release formats are protected

#### 2: AI-SUGGESTED NON-INFRINGING ALTERNATIVE

Feature	Al Suggestion
Proposed Compound	Huperzine A (a natural acetylcholinesterase inhibitor)
Delivery Form	Sublingual tablet, fast-dissolving
Half-life	4-6 hours (shorter than Rivastigmine)
Justification	No overlap in formulation or mechanism of delivery. Natural origin excludes it from synthetic patent class. Available open-access in various regions.
Regulatory Precedent	GRAS status in China, clinical trial support in EU

### 3: FORMULATION DESIGN (Alternative Product)

Product Name (internal)

Dosage Form

250 mcg Huperzine A, sublingual tablet

Excipients

Mannitol, Crospovidone, Magnesium Stearate, Natural Citrus Flavor

Bioavailability

73% (simulated vs. 56% oral Rivastigmine)

Onset Time

Recommended Usage

Twice daily

Cost per Unit (estimated)

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### 4: AI EFFICACY SIMULATION (vs. Rivastigmine)

Criteria	Rivastigmine	Proposed Formulation
Acetylcholinesterase Inhibition	86%	79%
Cognitive Score Improvement (Simulated MMSE)	+2.7	+2.4
Half-life	10 hrs (SR)	4 hrs
Patient Compliance	71%	82% (due to faster action & taste masking)

## 5: LEGAL & RISK ANALYSIS

No shared compound claims
 No infringement of coating/process/polymers
 Legal clearance across India, South Africa, Brazil
 Pending review in EU due to herbal classification

**LEGAL COMPARISON CHART** 

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EFFICACY SIMULATION GRAPHS (EXCEL)

**FULL TECHNICAL REPORT (PDF)** 

CHEMICAL STRUCTURES (.MOL)

DRAFT DOSSIER FOR REGULATORY SUBMISSION

